This listing of the claims replaces any and all prior versions and listings of claims in the application:

LISTING OF THE CLAIMS

1. (Currently amended) A composition comprising a biologically active compound, a transport moiety and a self-immolating linker moiety linking the biologically active compound and the transport moiety, wherein the transport moiety comprises a structure selected from the group consisting of $(ZY)_nZ$, $(ZYZ)_nZ$, $(ZYY)_nZ$, and $(ZYYY)_nZ$, wherein each Z is L-arginine or D-arginine, and each Y is independently an amino acid that does not comprise an amidino or guanidino moiety, and wherein n is an integer of from 2 to 10, wherein the conjugate has a structure selected from

$$R^{1}$$
— X — $(CH_{2})_{k}$ — A — C — $(CH_{2})_{m}$ — N — $(CH_{2})_{n1}$ — Q — R^{3}

$$R^{1}$$
— X — $(CH_{2})_{k}$ — R^{4} — $(CH_{2})_{m}$ — C — Q — R^{3}

$$R^{1}$$
— X — $(CH_{2})_{k}$ — C — Q — R^{3}

wherein

 R^1 is the biologically active compound;

X is a linkage between a functional group on the biologically active compound and a functional group on the linker between R^1 and R^3 ;

Q is a linkage between a functional group on the transport moiety and a functional group on the linker between R^1 and R^3 ;

A is N or CH;

Ar is a substituted or unsubstituted aryl group, wherein the methylene and oxygen substituents are either *ortho* or *para* to one another;

R² is hydrogen, alkyl, aryl, arylalkyl, acyl or allyl;

 R^3 is the transport moiety;

 R^4 is S, O, NR^6 or CR^7R^8 ;

 R^{4a} is S, O, NR^6 or $CR^{7a}R^{8a}$;

R⁵ is OH, SH, NHR⁶, or -CONH₂;

R^{5a} is H, OH, SH, NHR⁶, or -CONH₂;

R⁶ is hydrogen, alkyl, aryl, arylalkyl, acyl or allyl;

R⁷ and R⁸ are independently hydrogen, alkyl or arylalkyl; and

R^{7a} and R^{8a} are independently hydrogen or alkyl; and

k and m are independently either 1 or 2; and

n1 is an integer of from 1 to 10.

- 2. (original) The composition according to claim 1, wherein each Y is independently selected from the group consisting of alanine, cysteine, aspartic acid, glutamic acid, phenylalanine, glycine, histidine, isoleucine, lysine, leucine, methionine, asparagine, proline, glutamine, serine, threonine, valine, tryptophan, hydroxyproline, tyrosine, γ-amino butyric acid, β-alanine, sarcosine and ε-amino caproic acid.
- 3. (Original) The composition according to claim 1, wherein the transport moiety comprises the structure $(ZYZ)_nZ$, and wherein n is an integer ranging from 2 to 5.
- 4. (previously presented) The composition according to claim 1, wherein the transport moiety comprises the structure $(ZY)_nZ$ and wherein n is an integer ranging from 4 to 10.
- 5. (Original) The composition according to claim 1, wherein the transport moiety comprises the structure (ZYY)_nZ, and wherein n is an integer ranging from 4 to 10.

6. (Original) The composition according to claim 1, wherein the transport moiety comprises the structure (ZYYY)_nZ, and wherein n is an integer ranging from 4 to 10.

7. Canceled

- 8. (original) The composition according to claim 1, wherein Y is a gene-encoded amino acid.
- 9. (Original) The composition according to claim 1, wherein Y is an amino acid other than a gene-encoded amino acid.
- 10. (Original) The composition according to claim 3, wherein each Y is independently selected from the group consisting of glycine, γ -amino butyric acid, β -alanine and ϵ -amino caproic acid, and n is 3 or 4.
 - 11. (Currently amended) A composition, comprising:
- a biologically active compound, a transport moiety and a self-immolating linker moiety linking the biologically active compound and the transport moiety,

wherein the transport moiety comprises a structure <u>selected from the group consisting</u> of $(ZY)_nZ$, $(ZYZ)_nZ$, $(ZYY)_nZ$, and $(ZYYY)_nZ$,

each Z is L-arginine or D-arginine, each Y is independently glycine, γ -amino butyric acid, β -alanine or ϵ -amino caproic acid, and n is 6, 7 or 8, wherein the conjugate has a structure selected from

$$\frac{R^{1}-X-(CH_{2})_{k}-\overset{R^{5a}}{\overset{L}{C}-Q-R^{3}}_{H}_{, and}}{O_{, and}}$$

$$R^{1}-X-CH_{2}-Ar-O-\overset{H}{C}-(CH_{2})_{k}-R^{4a}-(CH_{2})_{m}-\overset{R^{5a}}{\overset{L}{C}-Q-R^{3}}_{H}$$

 R^1 is the biologically active compound;

X is a linkage between a functional group on the biologically active compound and a functional group on the linker between R^1 and R^3 ;

Q is a linkage between a functional group on the transport moiety and a functional group on the linker between R^1 and R^3 ;

A is N or CH;

Ar is a substituted or unsubstituted aryl group, wherein the methylene and oxygen substituents are either *ortho* or *para* to one another;

R² is hydrogen, alkyl, aryl, arylalkyl, acyl or allyl;

R³ is the transport moiety;

 $\underline{R^4}$ is S, O, NR^6 or CR^7R^8 ;

 R^{4a} is S, O, NR^6 or $CR^{7a}R^{8a}$;

R⁵ is OH, SH, NHR⁶, or -CONH₂;

R^{5a} is H, OH, SH, NHR⁶, or -CONH₂;

R⁶ is hydrogen, alkyl, aryl, arylalkyl, acyl or allyl;

R⁷ and R⁸ are independently hydrogen, alkyl or arylalkyl; and

R^{7a} and R^{8a} are independently hydrogen or alkyl; and

k and m are independently either 1 or 2; and

n1 is an integer of from 1 to 10.

12. (Original) The composition according to claim 5, wherein each Y is independently selected from the group consisting of glycine, γ -amino butyric acid, β -alanine and ϵ -amino caproic acid, and n is 6, 7 or 8.

- 13. (Original) The composition according to claim 6, wherein each Y is independently selected from the group consisting of glycine, γ -amino butyric acid, β -alanine and ϵ -amino caproic acid, and n is 6, 7 or 8.
- 14. (Currently amended) The composition according to claim 1, wherein the conjugate has the following structure:

R¹ is the biologically active compound;

X is a linkage between a functional group on the biologically active compound and a functional group on the linker between R¹ and R²;

Q is a linkage between a functional group on the transport moiety and a functional group on the linker between R¹ and R³;

A is N or CH;

R²-is hydrogen, alkyl, aryl, arylalkyl, acyl or allyl;

R³ is the transport moiety;

k and m are independently either 1 or 2; and

n is an integer of from 1 to 10.

15. (Original) The composition according to claim 14, wherein each of X and Q is independently selected from the group consisting of -C(O)O-, -O-C(O)-, -C(O)NH-, -NH-C(O)-, -OC(O)NH-, -S-S-, -C(S)O-, -C(S)NH-, -NHC(O)NH-, -SO₂NH-, -SONH-, phosphate, phosphonate and phosphinate.

- 16. (Original) The composition according to claim 14, wherein each of X and Q is independently selected from the group consisting of -C(O)O-, -O-C(O)-, -C(O)NH-, -NH-C(O)-, -OC(O)NH- and -NHC(O)NH-.
- 17. (Currently amended) The composition according to claim 1, wherein the conjugate has the following structure:

$$R^{1}$$
— X — $(CH_{2})_{k}$ — R^{4} — $(CH_{2})_{m}$ — C — C — Q — R^{3}

R¹ is the biologically active compound;

X is a linkage between a functional group on the biologically active compound and a functional group on the linker between R⁴ and R³;

Q is a linkage between a functional group on the transport moiety and a functional group on the linker between R⁺ and R³;

R³ is the transport moiety;

R⁴ is S, O, NR⁶ or CR⁷R⁸;

R⁵ is OH, SH₂ NHR⁶, or -CONH₂;

R⁶ is hydrogen, alkyl, aryl, arylalkyl, acyl or allyl;

R⁷ and R⁸ are independently hydrogen, alkyl or arylalkyl; and

k and m are independently either 1 or 2.

- 18. (Original) The composition according to claim 17 wherein each of X and Q is independently selected from the group consisting of -C(O)O-, -O-C(O)-, -C(O)NH-, -NH-C(O)-, -OC(O)NH-, -S-S-, -C(S)O-, -C(S)NH-, -NHC(O)NH-, -SO₂NH-, -SONH-, phosphate, phosphonate and phosphinate.
- 19. (Original) The composition according to claim 17, wherein each of X and Q is independently selected from the group consisting of -C(O)O-, -O-C(O)-, -C(O)NH-, -NH-C(O)-, -OC(O)NH- and -NHC(O)NH-.

20. (Currently amended) The composition according to claim 1, wherein the conjugate has the following structure:

$$\frac{R^{5}}{R^{+}X-(CH_{2})_{k}-C-Q-R^{3}}$$

$$R^{1}-X-(CH_{2})_{k}-C-Q-R^{3}$$

wherein:

R¹ is the biologically active compound;

X is a linkage between a functional group on the biologically active compound and a functional group on the linker between R¹ and R³;

Q is a linkage between a functional group on the transport moiety and a functional group on the linker between R¹ and R²;

R³ is the transport moiety;

R⁵ is H, OH, SH, NHR⁶, or -CONH₂;

R⁶ is hydrogen, alkyl, aryl, arylalkyl, acyl or allyl; and k is 1 or 2.

- 21. (Original) The composition according to claim 20, wherein each of X and Q is independently selected from the group consisting of -C(O)O-, -O-C(O)-, -C(O)NH-, -NH-C(O)-, -OC(O)NH-, -S-S-, -C(S)O-, -C(S)NH-, -NHC(O)NH-, -SO₂NH-, -SONH-, phosphate, phosphonate and phosphinate.
- 22. (Original) The composition according to claim 20, wherein each of X and Q is independently selected from the group consisting of -C(O)O-, -O-C(O)-, -C(O)NH-, -NH-C(O)-, -OC(O)NH- and -NHC(O)NH-.
- 23. (Currently amended) The composition according to claim 1, wherein the conjugate has the following structure:

$$\begin{array}{c} O & R^{5} \\ \hline R^{1}-X-CH_{2}-Ar-O-C-(CH_{2})_{k}-R^{4}-(CH_{2})_{m}-C-Q-R^{3} \\ \hline \\ R^{1}-X-CH_{2}-Ar-O-C-(CH_{2})_{k}-R^{4a}-(CH_{2})_{m}-C-Q-R^{3} \\ \hline \end{array}$$

R¹ is the biologically active compound;

X is a linkage between a functional group on the biologically active compound and a functional group on the linker between R⁴ and R³;

Q is a linkage between a functional group on the transport moiety and a functional group on the linker between R¹ and R³:

Ar is a substituted or unsubstituted aryl group, wherein the methylene and oxygen substituents are either *ortho* or *para* to one another;

R³ is the transport moiety;

R⁴ is S, O, NR⁶ or CR⁷R⁸;

R⁵ is H, OH, SH, CONHR⁶ or NHR⁶;

R⁶ is hydrogen, alkyl, aryl, arylalkyl, acyl or allyl;

R⁷ and R⁸ are independently hydrogen or alkyl; and,

k and m are independently either 1 or 2.

- 24. (Original) The composition according to claim 23, wherein each of X and Q is independently selected from the group consisting of -C(O)O-, -O-C(O)-, -C(O)NH-, -NH-C(O)-, -OC(O)NH-, -S-S-, -C(S)O-, -C(S)NH-, -NHC(O)NH-, -SO₂NH-, -SONH-, phosphate, phosphonate and phosphinate.
- 25. (Original) The composition according to claim 23, wherein each of X and Q is independently selected from the group consisting of -C(O)O-, -O-C(O)-, -C(O)NH-, -NH-C(O)-, -OC(O)NH- and -NHC(O)NH-.
- 26. (Currently amended) The composition according to claim 16, wherein A is N, R^2 is benzyl, k, m and [[n]] n1 are 1, and X is -OC(O)-.

27-29. (cancelled)

30. (Currently amended) A method for increasing the transport of a biologically active compound across a biological membrane comprising:

administering a composition according to claim 1 comprising a biologically active compound, a transport moiety, and a linker capable of self-immolation linking the biologically active compound and the transport moiety, wherein the transport compound comprises a structure selected from the group consisting of (ZYZ)_nZ, (ZY)_nZ, (ZYY)_nZ and (ZYYY)_nZ, wherein Z is L arginine or D arginine, and wherein Y is an amino acid that does not comprise an amidino or guanidino moiety, and wherein n is an integer ranging from 2 to 10,

wherein transport of the biologically active biologically active compound across the biological membrane is increased relative to transport of the biologically active compound in the absence of said transport moiety.

- 31. (Canceled).
- 32. (Currently amended) The method of claim [[31]] <u>30</u>, wherein the conjugate has the following structure:

wherein:

R¹ is the biologically active compound;

X is a linkage between a functional group on the biologically active compound and a functional group on the linker between R^4 and R^3 ;

Q is a linkage between a functional group on the transport moiety and a functional group on the linker between R¹ and R³;

A is N or CH;

R² is hydrogen, alkyl, aryl, arylalkyl, acyl or allyl;

R³ is a transport moiety;

k and m are independently either 1 or 2; and

n is an integer of from 1 to 10.

33. (Currently amended) The method of claim [[31]] <u>30</u>, wherein the conjugate has the following structure:

$$R^{1}$$
— X — $(CH_{2})_{k}$ — R^{4} — $(CH_{2})_{m}$ — C — C — Q — R^{3}

wherein:

R¹ is the biologically active compound;

X is a linkage between a functional group on the biologically active compound and a functional group on the linker between R⁴ and R³;

Q is a linkage between a functional group on the transport moiety and a functional group on the linker between R¹ and R³;

R³ is a transport moiety;

R⁴ is S, O, NR⁶ or CR⁷R⁸;

R⁵ is OH, SH, NHR⁶, or CONH₂;

R⁶ is hydrogen, alkyl, aryl, arylalkyl, acyl or allyl;

R⁷-and R⁸-are independently hydrogen, alkyl or arylalkyl; and

k and m are independently either 1 or 2.

34. (Currently amended) The method of claim [[31]] <u>30</u>, wherein the conjugate has the following structure:

$$\begin{array}{c}
R^{5} \\
\hline
R^{+}X-(CH_{2})_{k}-C-Q-R^{3} \\
H
\end{array}$$

R⁴ is the biologically active compound;

X is a linkage between a functional group on the biologically active compound and a functional group on the linker between R¹ and R³;

Q is a linkage between a functional group on the transport moiety and a functional group on the linker between R¹ and R²:

R³ is the transport moiety;

R⁵ is H, OH, SH, NHR⁶, or -CONH₂;

R⁶ is hydrogen, alkyl, aryl, arylalkyl, acyl or allyl; and

k is 1 or 2.

35. (Currently amended) The method of claim [[31]] <u>30</u>, wherein the conjugate is of the following structure:

$$\begin{array}{c} O & R^{5} \\ \hline R^{1}-X-CH_{2}-Ar-O-C-(CH_{2})_{k}-R^{4}-(CH_{2})_{m}-C-Q-R^{3} \\ \\ O & R^{5a} \\ R^{1}-X-CH_{2}-Ar-O-C-(CH_{2})_{k}-R^{4a}-(CH_{2})_{m}-C-Q-R^{3} \\ \\ \end{array}$$

wherein:

R¹ is the biologically active compound;

X is a linkage between a functional group on the biologically active compound and a functional group on the linker between R^4 and R^3 ;

Q is a linkage between a functional group on the transport moiety and a functional group on the linker between R¹ and R³;

Ar is a substituted or unsubstituted aryl group, wherein the methylene and oxygen substituents are either *ortho* or *para* to one another;

R³ is the transport moiety;

R⁴-is S, O, NR⁶-or CR⁷R⁸;

R⁵-is H, OH, SH, CONHR⁶-or NHR⁶;

R⁶-is hydrogen, alkyl, aryl, arylalkyl, acyl or allyl;

R⁷-and R⁸-are independently hydrogen or alkyl; and,
k and m are independently either 1 or 2.

- 36. (Original) The composition of claim 1, wherein said linker moiety covalently links the biologically active compound and the transport moiety.
- 37. (Original) The composition of claim 1, wherein said linker moiety capable of self-immolation is configured so as to undergo intramolecular cleavage.
- 38. (Original) The composition of claim 1, wherein said linker moiety comprises a half-life in the range of between about 10 minutes and about 24 hours in water at 37 °C and at a pH of approximately 7.4.
 - 39. (Canceled).
 - 40. (Canceled).